

## ESTER DERIVATIVES

Publication number: WO0204402

Publication date: 2002-01-17

Inventor: OGINO YOSHIO (JP); KURIHARA HIDEKI (JP); MATSUDA KENJI (JP); NUMAZAWA TOMOSHIGE (JP); OTAKE NORIKAZU (JP); NOGUCHI KAZUHITO (JP)

Applicant: BANYU PHARMA CO LTD (JP); OGINO YOSHIO (JP); KURIHARA HIDEKI (JP); MATSUDA KENJI (JP); NUMAZAWA TOMOSHIGE (JP); OTAKE NORIKAZU (JP); NOGUCHI KAZUHITO (JP)

Classification:  
- international: A61P11/00; A61P11/06; A61P27/16; A61P43/00; C07C219/10; C07C219/14; C07D205/04; C07D207/08; C07D207/12; C07D209/52; C07D211/22; C07D211/42; C07D211/46; C07D211/70; C07D239/06; C07D451/02; C07D451/06; C07D471/08; C07D471/10; C07D487/10; C07D498/10; A61P11/00; A61P27/00; A61P43/00; C07C219/00; C07D205/00; C07D207/00; C07D209/00; C07D211/00; C07D239/00; C07D451/00; C07D471/00; C07D487/00; C07D498/00; (IPC1-7): C07C219/10; A61K31/222; A61K31/395; A61K31/397; A61K31/40; A61K31/403; A61K31/407; A61K31/435; A61K31/438; A61K31/439; A61K31/4409; A61K31/452; A61K31/4525; A61K31/5386; A61K31/55; A61P11/00; A61P11/06; A61P27/16; A61P43/00; C07C219/22; C07C219/24; C07C251/08; C07C251/18; C07D205/04; C07D207/08; C07D207/12; C07D209/52; C07D211/22; C07D211/46; C07D211/70; C07D221/24; C07D239/06; C07D295/125; C07D405/12; C07D451/02; C07D471/10; C07D487/10; C07D498/10

- european: C07C219/10; C07C219/14; C07D205/04; C07D207/08A; C07D207/12; C07D209/52; C07D211/22; C07D211/42; C07D211/46; C07D211/70; C07D239/06B3; C07D239/06C; C07D451/02B; C07D451/06D; C07D471/08; C07D471/10; C07D487/10; C07D498/10

Application number: WO2001JP05987 20010710

Priority number(s): JP20000210591 20000711

## Also published as:

EP1302458 (A1)  
US6846835 (B2)  
US2003191316 (A1)  
CA2415468 (A1)

attached

## Cited documents:

WO9821183  
EP0140434  
FR1352332[Report a data error here](#)

## Abstract of WO0204402

Compounds of the general formula (1), which exhibit selective muscarinic M3 receptor antagonism, little have side effects, and are suitable for administration by inhalation and useful as therapeutic agents for respiratory system diseases or the like: (1) wherein A is a group of the general formula (a0) or (b0): (a0) (b0) Ar is aryl or heteroaryl, any of which may be substituted; B<1> and B<2> are each an aliphatic hydrocarbon group; R<1> is fluorinated cycloalkyl; R<2>, R<3> and R<4> are each lower alkyl, or a single bond or alkylene, any of which is bonded to B<1>, or alternatively R<2> and R<3> may be united to form alkylene; R<5> and R<7> are each hydrogen, lower alkyl, or a single bond or alkylene, any of which is bonded to B<2>; R<6> is hydrogen, lower alkyl, or N(R<8>)R<9>; and X<-> is an anion.

Compounds of the general formula (1), which exhibit selective muscarinic M3 receptor antagonism, little have side effects, and are suitable for administration by inhalation and useful as therapeutic agents for respiratory system diseases or the like: (1) wherein A is a group of the general formula (a0) or (b0): (a0) (b0) Ar is aryl or heteroaryl, any of which may be substituted; B<1> and B<2> are each an aliphatic hydrocarbon group; R<1> is fluorinated cycloalkyl; R<2>, R<3> and R<4> are each lower alkyl, or a single bond or alkylene, any of which is bonded to B<1>, or alternatively R<2> and R<3> may be united to form alkylene; R<5> and R<7> are each hydrogen, lower alkyl, or a single bond or alkylene, any of which is

bonded to B<2>; R<6> is hydrogen, lower alkyl, or N(R<8>)R<9>; and X<-> is an anion.

---

Data supplied from the **esp@cenet** database - Worldwide

